



Automated Analysis of Natural Speech in Amyotrophic Lateral Sclerosis

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Study funding:

National Institutes of Health (AG017586, AG053940, AG052943, NS088341, DC013063, AG054519), the Institute on Aging at the University of Pennsylvania, the Alzheimer's Association (AACSF-18-567131), an anonymous donor, and the Wyncote Foundation.

Disclosures:

Dr. Grossman participates in clinical trials sponsored by Alector, Eisai and Biogen that are unrelated to this study. He receives research support from NIH, Biogen and Avid that is unrelated to this study. He receives financial support from Neurology for his work as an Associate Editor. C.M receives research funding from Biogen, Inc and provides consulting services for Invicro and Axon Advisors on behalf of Translational Bioinformatics, LLC. He also receives an honorarium as Associate Editor of NeuroImage: Clinical. All other authors have nothing to disclose.

Background: What do we know about speech in ALS?

- Impairments in
 - grammatical processing
 - action verb knowledge
 - discourse and social communication
 - Reading time (prolonged)

Speech is a multidimensional skill requiring the collaboration of multiple cognitive and motor domains. Our study assesses the interactions of motor and cognitive impairments on acoustic-prosodic aspects of speech in ALS and ALS-FTD.

Potential motor & cognitive effects on speech in ALS

Cognitive

- Behavioral (more common)
 - Social
 - Executive
- Linguistic
 - Agrammatism (more common)
 - Semantic
 - Audio-verbal

Motor

- Bulbar disease
- Reduced respiratory capacity

Study objectives

- Characterize acoustic-prosodic properties of speech in ALS spectrum
- Identify motor vs. cognitive effects on speech in ALS spectrum
- Implementation and validation of automated speech recognition in clinical settings

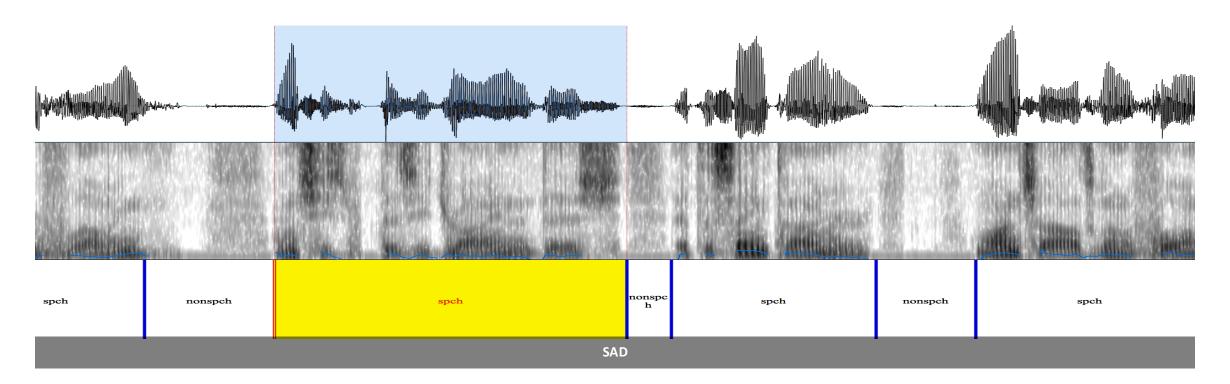
Hypothesis

• Acoustic features of speech in speakers with ALS and ALS-FTD can provide distinct markers that reflect motor and cognitive impairments.



Methods

- Digitized narrative speech samples picture description task
- Automatic segmentation with a speech activity detector (SAD)



Methods – patient groups

Clinical & Demographic characteristics - Mean (SD)						
	ALS	ALS-FTD	HC	bvFTD	naPPA	р
n	44	23	33	90	23	
Sex = Male (%)	23 (52.3)	17 (73.9)	13 (39.4)	56 (62.2)	12 (52.2)	0.076
Age (y)	62.4 (10.4)	64.6 (8.8)	67.6 (6.2)	62.8 (8.7)	70.6 (9.2)	<0.001
Education (y, n=210)	15.1 (2.8)	13.4 (2.1)	16.1 (2.5)	15.8 (2.8)	15.2 (3.0)	0.004
Symptom duration (y, n=179)	3.7 (2.6)	4.4 (3.6)	NA	4.4 (3.2)	3.4 (1.8)	0.408
Bulbar disease = yes (%)	16 (37.2)	8 (34.8)	NA	NA	NA	NA
%FVC (n=61)	78.5 (28.1)	63.7 (23.5)	NA	NA	NA	0.042
ALSFRS-R total score (0-48, n=61)	35.0 (7.5)	34.2 (7.1)	NA	NA	NA	0.709
ECAS total score (0-136, n=53)	115.1 (5.2)	84.7 (19.1)	NA	NA	NA	<0.001

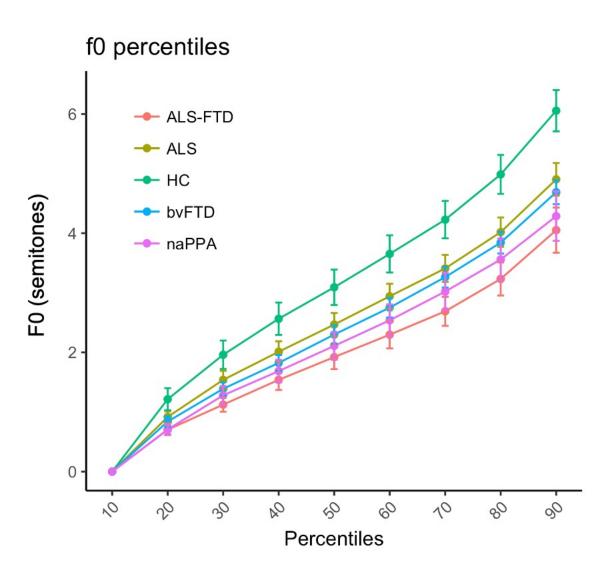
ALS - Amyotrophic lateral sclerosis; ALSFRS-R - ALS functional rating scale revised; ALS-FTD - ALS with frontotemporal dementia; bvFTD - behavioral variant FTD; ECAS - Edinburgh Cognitive Assessment Scale; %FVC - forced vital capacity (% of predicted by age); HC - healthy control; MMSE - minimental status examination; NA - not available; naPPA - nonfluent/agrammatic variant of Primary progressive aphasia; SD - standard deviation; y - years.

Methods – cont.

- Pitch tracking of continuous speech segments
- Duration measures for speech and silent pause segments
- Calculated acoustic measures: fundamental frequency (f0) range, mean speech and total speech durations, pause rate.
- Statistical analyses:
 - Group comparisons (controlling for age and education, adjusting for multiple comparisons)
 - Linear regression models:
 - acoustic measure ~ cognitive test score + motor function
 - cortical atrophy (MRIT1) ~ acoustic measure

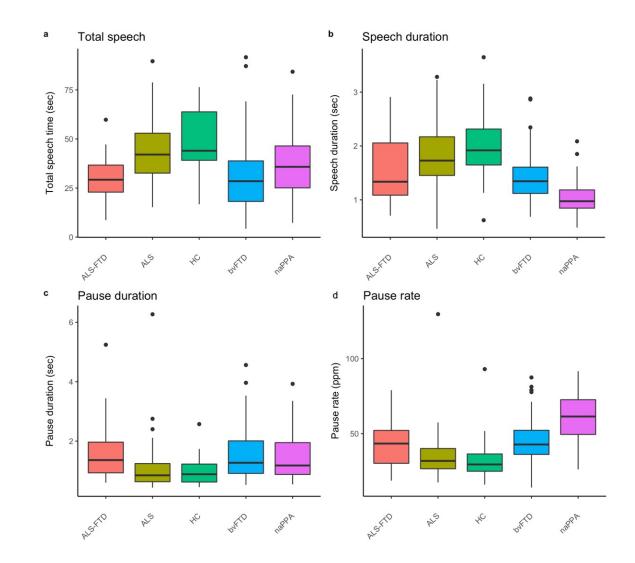
Results

 f0 range is narrow in ALS spectrum disorders compared with normal speakers



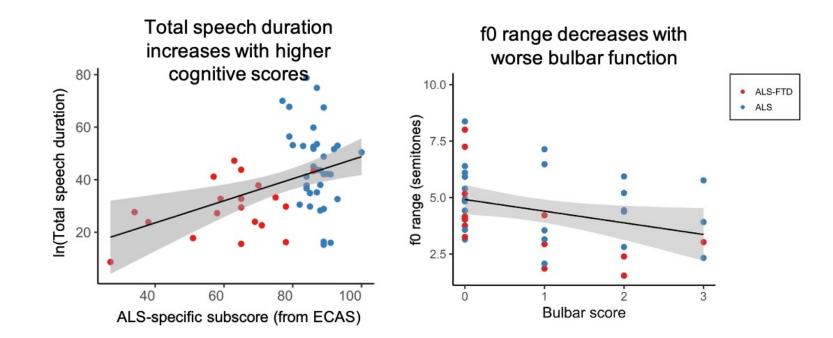
Results - cont.

- Mean speech segment duration and total speech duration were shorter in ALS-FTD compared with normal and ALS speakers
- Pause duration and rate were impaired in ALS-FTD compared with normal and ALS speakers
- ALS-FTD speakers' durational acoustic features most resemble bvFTD speakers'



Results – Clinical correlates

- Impaired f0 range was related to bulbar disease (beta=-0.59, p=0.012)
- Speech duration (beta=0.38, p=0.006) was related to cognitive impairment independent of respiratory function



Results – Anatomical correlates

- Impaired f0 range (red) was associated with atrophy in primary motor cortex and left peri-Sylvian regions.
- Total speech duration (blue) was associated with atrophy in the IFG bilaterally.
- Lt. frontal operculum (magenta) linked to both f0 and speech



Conclusions

- Speech samples in ALS spectrum disorders can provide highly objective and reproducible markers of disease derived purely from the acoustic signal.
- Acoustic markers relate to prosodic elements of natural language such as fluency and intonation and reflect specific motor and cognitive impairments in ALS.

A full-length manuscript of this study was accepted for publication in Neurology.

Resources

Please refer questions and comments to the corresponding author:

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Acknowledgements

Penn FTDC

- Katya Rascovsky
- RA list
- Sanjana Shellikeri
- Natalia Parjane
- Our patients and their caregivers

Penn LDC

- Chris Cieri
- Catelin Cieri
- SAD writer
- Programmers / transcribers